SENATE BILL No. 258

DIGEST OF INTRODUCED BILL

Citations Affected: IC 15-15-13; IC 16-18-2; IC 16-42-28; IC 21-45-7; IC 34-30-2-83.9; IC 35-48-4.

Synopsis: Cannabidiol for the treatment of epilepsy. Provides that certain prohibitions against granting a license to a grower or handler of industrial hemp do not apply to growers and manufacturers that process cannabidiol (CBD) and meet certain requirements. Requires the state seed commissioner to establish testing standards for CBD. Establishes requirements for facilities and manufacturers that manufacture or process CBD. Allows CBD that is manufactured and tested in Indiana to be used by certain physicians, patients, and caregivers. Establishes a CBD registry for certain physicians, patients, and caregivers for the use of CBD from hemp in the treatment of a child with intractable epilepsy. Establishes a pilot study registry for physicians interested in studying the use of CBD in the treatment of intractable epilepsy. Requires the state department of health to develop and maintain both registries. Provides civil, criminal, and administrative immunity for: (1) physicians in the use of CBD in the treatment of intractable epilepsy; (2) certain growers of industrial hemp; and (3) facilities and manufacturers of CBD; if certain requirements are met. Exempts caregivers and patients from criminal penalties for possession or use of CBD if the caregivers and patients are registered with the state department and are using the CBD for the patient and in the manner approved for registration. Encourages state educational institutions to research the use of CBD in the treatment of intractable epilepsy.

Effective: July 1, 2016.

Tomes

January 7, 2016, read first time and referred to Committee on Health & Provider Services.



Second Regular Session 119th General Assembly (2016)

PRINTING CODE. Amendments: Whenever an existing statute (or a section of the Indiana Constitution) is being amended, the text of the existing provision will appear in this style type, additions will appear in this style type, and deletions will appear in this style type.

Additions: Whenever a new statutory provision is being enacted (or a new constitutional provision adopted), the text of the new provision will appear in **this style type**. Also, the word **NEW** will appear in that style type in the introductory clause of each SECTION that adds a new provision to the Indiana Code or the Indiana Constitution.

Conflict reconciliation: Text in a statute in *this style type* or *this style type* reconciles conflicts between statutes enacted by the 2015 Regular Session of the General Assembly.

SENATE BILL No. 258

A BILL FOR AN ACT to amend the Indiana Code concerning health.

Be it enacted by the General Assembly of the State of Indiana:

1	SECTION 1. IC 15-15-13-2.5 IS ADDED TO THE INDIANA
2	CODE AS A NEW SECTION TO READ AS FOLLOWS
3	[EFFECTIVE JULY 1, 2016]: Sec. 2.5. As used in this chapter
4	"cannabidiol" has the meaning set forth in IC 16-42-28-1.
5	SECTION 2. IC 15-15-13-5, AS ADDED BY P.L.165-2014
6	SECTION 1, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
7	JULY 1, 2016]: Sec. 5. As used in this chapter, "handler" means:
8	(1) an individual, a partnership, a company, or a corporation tha
9	receives industrial hemp for scientific research, or for processing
0	into commodities, products, or agricultural hemp seed; or
1	(2) a facility or manufacturer that manufactures or processes
2	cannabidiol.
3	SECTION 3. IC 15-15-13-15, AS ADDED BY P.L.165-2014
4	SECTION 1, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
5	JULY 1, 2016]: Sec. 15. (a) This section does not apply to a grower
6	or handler who grows, supplies, or processes industrial hemp into
7	cannabidiol under IC 15-15-13-18.



1	(a) (b) Notwithstanding any other law, the state seed commissioner
2	may not grant any license until the state seed commissioner has secured
3	any necessary permissions, waivers, or other form of legal status by the
4	United States Drug Enforcement Agency or other appropriate federal
5	agency concerning industrial hemp.
6	(b) (c) The state seed commissioner shall apply for any necessary
7	permissions, waivers, or other forms of legal status by the United States
8	Drug Enforcement Agency or other appropriate federal agency that are
9	necessary to implement this chapter. before January 1, 2015.
10	(c) (d) The state seed commissioner may not implement a waiver
11	under this section until the state seed commissioner files an affidavit
12	with the governor attesting that the federal permission or waiver
13	applied for under this section is in effect. The state seed commissioner
14	shall file the affidavit under this subsection not later than five (5) days
15	after the state seed commissioner is notified that the waiver is
16	approved.
17	(d) (e) If the state seed commissioner receives a waiver permission
18	under this section from all the appropriate federal agencies and the
19	governor receives the affidavit filed under subsection (c), (d), the state
20	seed commissioner shall implement this chapter, subject to the terms
21	and conditions of the permission or waiver received, not more than
22	sixty (60) days after the governor receives the affidavit.
23	SECTION 4. IC 15-15-13-18 IS ADDED TO THE INDIANA
24	CODE AS A NEW SECTION TO READ AS FOLLOWS
25	[EFFECTIVE JULY 1, 2016]: Sec. 18. (a) This section does not apply
26	to a facility or manufacturer that is:
27	(1) a pharmaceutical manufacturer operating under the
28	approval of the federal Food and Drug Administration; or
29	(2) a state educational institution.
30	(b) The state seed commissioner shall adopt rules to establish
31	testing standards for cannabidiol that govern the analyzing and
32	testing of cannabidiol (CBD) for purity, potency, and
33	tetrahydrocannabinol (THC) levels.
34	(c) A facility or manufacturer that manufacturers or processes
35	cannabidiol in Indiana shall do the following:
36	(1) Submit a research plan to the state seed commissioner
37	concerning the investigation of:
38	(A) the processing of industrial hemp;
39	(B) the use of equipment or other methods to extract
40	cannabidiol from industrial hemp;
41	(C) the extraction of cannabidiol from different varieties
42	of industrial hemp; or



1	(D) other legitimate research concerning industrial hemp.
2	(2) Comply with the testing standards established under
3	subsection (b).
4	(3) Affix a copy of the test results on each container that
5	contains cannabidiol.
6	(4) Maintain sanitary conditions in the areas where
7	cannabidiol is manufactured or processed.
8	(5) Maintain adequate security over manufactured or
9	processed cannabidiol.
10	(6) Supply cannabidiol only to:
11	(A) physicians;
12	(B) patients; or
13	(C) caregivers;
14	under IC 16-42-28.
15	(d) A facility or manufacturer that manufacturers or processes
16	cannabidiol and meets the requirements under this chapter shall
17	be licensed as a handler under this chapter.
18	(e) A grower may grow and supply industrial hemp to a facility
19	or manufacturer that manufacturers or processes cannabidiol
20	under this section.
21	(f) The state seed commissioner may inspect a facility or
22	manufacturer that manufacturers or processes cannabidiol to
23	ensure compliance with this section.
24	SECTION 5. IC 16-18-2-43.5 IS ADDED TO THE INDIANA
25	CODE AS A NEW SECTION TO READ AS FOLLOWS
26	[EFFECTIVE JULY 1, 2016]: Sec. 43.5. "Cannabidiol", for purposes
27	of IC 16-42-28, has the meaning set forth in IC 16-42-28-1.
28	SECTION 6. IC 16-18-2-48.7 IS ADDED TO THE INDIANA
29	CODE AS A NEW SECTION TO READ AS FOLLOWS
30	[EFFECTIVE JULY 1, 2016]: Sec. 48.7. "Caregiver", for purposes
31	of IC 16-42-28, has the meaning set forth in IC 16-42-28-2.
32	SECTION 7. IC 16-18-2-191.7 IS ADDED TO THE INDIANA
33	CODE AS A NEW SECTION TO READ AS FOLLOWS
34	[EFFECTIVE JULY 1, 2016]: Sec. 191.7. "Intractable epilepsy", for
35	purposes of IC 16-42-28, has the meaning set forth in
36	IC 16-42-28-3.
37	SECTION 8. IC 16-18-2-272 IS AMENDED TO READ AS
38	FOLLOWS [EFFECTIVE JULY 1, 2016]: Sec. 272. (a) "Patient", for
39	purposes of IC 16-27-1, has the meaning set forth in IC 16-27-1-6.
41	individual who has been accepted and assured care by a health facility.
40 41	(b) "Patient", for the purposes of IC 16-28 and IC 16-29, means an individual who has been accepted and assured care by a health facility.

(c) "Patient", for purposes of IC 16-36-1.5, has the meaning set forth



42

1	in IC 16-36-1.5-3.
2	(d) "Patient", for purposes of IC 16-39, means an individual who has
3	received health care services from a provider for the examination,
4	treatment, diagnosis, or prevention of a physical or mental condition.
5	(e) "Patient", for purposes of IC 16-42-28, has the meaning set
6	forth in IC 16-42-28-4.
7	SECTION 9. IC 16-18-2-282, AS AMENDED BY P.L.6-2012
8	SECTION 111, IS AMENDED TO READ AS FOLLOWS
9	[EFFECTIVE JULY 1, 2016]: Sec. 282. (a) "Physician", except as
10	provided in subsections (b) and (c), through (d), means a licensed
11	physician (as defined in section 202 of this chapter).
12	(b) "Physician", for purposes of IC 16-41-12, has the meaning set
13	forth in IC 16-41-12-7.
14	(c) "Physician", for purposes of IC 16-37-1-3.1 and IC 16-37-3-5.
15	means an individual who:
16	(1) was the physician last in attendance (as defined in section
17	282.2 of this chapter); or
18	(2) is licensed under IC 25-22.5.
19	(d) "Physician", for purposes of IC 16-42-28, has the meaning
20	set forth in IC 16-42-28-5.
21	(d) (e) "Physician", for purposes of IC 16-48-1, is subject to
22	IC 16-48-1-2.
23	SECTION 10. IC 16-42-28 IS ADDED TO THE INDIANA CODE
24	AS A NEW CHAPTER TO READ AS FOLLOWS [EFFECTIVE
25	JULY 1, 2016]:
26	Chapter 28. Drugs: Study and Use of Cannabidiol for the
27	Treatment of Epilepsy Program
28	Sec. 1. As used in this chapter, "cannabidiol" means the
29	cannabidiol (CBD) from a hemp plant that is manufactured or
30	processed by a:
31	(1) pharmaceutical manufacturer operating under the
32	approval of the federal Food and Drug Administration;
33	(2) state educational institution; or
34	(3) facility or manufacturer that is in compliance with the
35	requirements under IC 15-15-13-18.
36	Sec. 2. As used in this chapter, "caregiver" refers to a parent
37	legal guardian, health care representative, or custodian of a
38	patient.
39	Sec. 3. As used in this chapter, "intractable epilepsy" means a
40	seizure disorder that has been diagnosed by a physician in a patient
41	who has not responded to at least three (3) other seizure disorder
42	treatment options.



treatment options.

1	Sec. 4. As used in this chapter, "patient" refers to an individual
2	who is:
3	(1) less than eighteen (18) years of age; or
4	(2) at least eighteen (18) years of age but started treatment
5	with cannabidiol described under this chapter when the
6	individual was less than eighteen (18) years of age;
7	and who has been diagnosed with intractable epilepsy by a
8	physician.
9	Sec. 5. As used in this chapter, "physician" means an individual
10	who:
11	(1) is licensed under IC 25-22.5;
12	(2) is board certified in neurology; and
13	(3) is affiliated with a state educational institution.
14	Sec. 6. A physician who is registered and conducting a
15	registered pilot research study under this chapter may approve the
16	dispensing or use of cannabidiol to a registered caregiver or
17	registered patient for the treatment of intractable epilepsy.
18	Sec. 7. (a) The state department shall develop and implement a
19	cannabidiol registry for the registration of:
20	(1) physicians;
21	(2) patients; and
22	(3) caregivers;
23	for the use of cannabidiol in the treatment of patients with
24	intractable epilepsy.
25	(b) The cannabidiol registry must include a secure, electronic
26	online data base that is accessible by law enforcement agencies in
27	order to verify the registration of an individual.
28	(c) The state department shall register and issue a registration
29	card to a physician who:
30	(1) conducts a pilot research study;
31	(2) meets the requirements of this chapter;
32	(3) submits a completed registration form issued by the state
33	department; and
34	(4) pays the registration fee.
35	(d) The state department shall register and issue an individual
36	described in subsection (a)(2) or (a)(3) a registration card under
37	this section only if the individual meets the following requirements:
38	(1) The registrant under this subsection is:
39	(A) a caregiver who is at least eighteen (18) years of age; or
40	(B) a patient.
41	(2) The registrant under this subsection is an Indiana resident.
42	(3) The registrant under this subsection provides a certified



1	statement by a physician registered under this section that the
2	registrant is either the patient or caregiver of a patient who
3	meets the following requirements:
4	(A) The patient has been examined and diagnosed by the
5	physician to have intractable epilepsy.
6	(B) The patient has been recommended by the physician
7	for treatment by cannabidiol for the intractable epilepsy.
8	(C) The patient is to be included in the physician's pilot
9	research study registry under section 9 of this chapter.
0	(4) The registrant under this subsection submits a completed
11	registration application.
12	(5) The registrant under this subsection pays the registration
13	fee set by the state department.
14	(e) The state department shall develop the cannabidiol
15	registration application. The registration application for a
16	caregiver must ask for the following information:
17	(1) The caregiver's name and address.
18	(2) The patient's name and address.
19	(3) A copy of the caregiver's valid government issued photo
20	identification card.
21	(4) Any other relevant information the state department
22	considers necessary to implement this section.
23	(f) The state department shall charge each registrant a
24	registration fee of not more than fifty dollars (\$50) for an
25	individual's initial registration under this section to cover the costs
26	of implementing and administering the registry.
27	(g) Registration under this section is valid for one (1) year from
28	the date of issuance. The state department shall renew a
29	registration under this section for a registrant if the initial
30	registration is current or has been updated by the registrant and
31	the registrant continues to meet the registration requirements
32	under this chapter. The state department shall charge a renewal
33	fee of not more than twenty-five dollars (\$25).
34	Sec. 8. When a patient or caregiver registers for the cannabidiol
35	registry, the state department shall contact and provide the local
36	department of health where the patient or caregiver resides the
37	following information:
38	(1) The name and address of the patient or caregiver.
39	(2) Identifying information contained on the patient's or
10	caregiver's registration card.
1 1	(3) Any other information the state department determines is
12	necessary to disclose.



1	Sec. 9. (a) The state department shall establish and maintain a
2	pilot study registry for the monitoring of research performed by a
3	state educational institution as described in IC 21-45-7 concerning
4	the safety and efficacy of using cannabidiol in the treatment of
5	intractable epilepsy.
6	(b) A physician or the state educational institution seeking to
7	conduct research described in subsection (a) shall submit to the
8	state department an application to be included in the pilot study
9	registry that includes the following information:
10	(1) The name of the pilot study.
11	(2) The name of the affiliated state educational institution.
12	(3) The scientific and clinical parameters of the study.
13	(4) The protocols established to ensure patient safety.
14	(5) The name and address of any physician associated with the
15	pilot study.
16	(6) Any other information considered necessary by the state
17	department in order to determine the safety and evidence
18	based nature of the pilot study.
19	A physician may not conduct the research described in subsection
20	(a) until the pilot study has been certified by the state departmen
21	under subsection (c).
22	(c) The state department shall review each application
23	submitted under subsection (b) and include the pilot study on the
24	registry described in subsection (a) only after the state departmen
25	determines and certifies that the proposed pilot study does the
26	following:
27	(1) Adheres to minimum scientific methods.
28	(2) Protects patient safety.
29	(d) The state department may monitor a registered pilot study
30	under this section to ensure that the physician adheres to the
31	requirements set forth in subsection (c).
32	(e) A physician who has been approved for the pilot study
33	registry shall do the following while conducting the pilot study
34	research approved under this section:
35	(1) Maintain records of the evaluations and observations of a
36	patient participating in the pilot study, including the patient's
37	response to the cannabidiol treatment.
38	(2) Transmit the records described in subdivision (1) to the
39	state department upon the state department's request.
40	(f) The state department may charge a fee to cover
41	implementation and administration of the registry established



under this section.

8
Sec. 10. The state department shall maintain any medical records obtained under this chapter as confidential and the medical records may not be disclosed to the public. Sec. 11. (a) A physician registered under this chapter is immune
from civil, criminal, and administrative liability for approving, dispensing, or using cannabidiol in the treatment of a patient with
intractable epilepsy if the patient is validly registered under this chapter. (b) A grower under IC 15-15-13-18 is immune from civil, criminal, and administrative liability for growing or suppling

- (b) A grower under IC 15-15-13-18 is immune from civil, criminal, and administrative liability for growing or suppling industrial hemp to a manufacturer or grower under IC 15-15-13-18.
- (c) A facility or manufacturer under IC 15-15-13-18 is immune from civil, criminal, and administrative liability for manufacturing, processing, or suppling cannabidiol under IC 15-15-13-18 or this chapter.
- Sec. 12. (a) The state department shall study whether the registration under section 7 of this chapter should, based on sound medical findings, be extended to individuals who:
 - (1) are at least eighteen (18) years of age;
 - (2) have been examined and diagnosed by a physician to have intractable epilepsy; and
 - (3) have been recommended by the physician for treatment by cannabidiol for the intractable epilepsy.
- (b) Before November 1, 2017, the state department shall provide its findings under subsection (a) to the general assembly in an electronic format under IC 5-14-6. This subsection expires December 31, 2017.
- Sec. 13. The state department may adopt rules under IC 4-22-2 necessary to implement this chapter.
- SECTION 11. IC 21-45-7 IS ADDED TO THE INDIANA CODE AS A **NEW** CHAPTER TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2016]:
- Chapter 7. Research on Cannabidiol for Treatment of Epilepsy Sec. 1. As used in this chapter, "cannabidiol" means the cannabidiol from a hemp plant.
- Sec. 2. As used in this chapter, "intractable epilepsy" means a seizure disorder that has been diagnosed by a physician (as defined in IC 16-42-28-5) in a patient who has not responded to at least three (3) other seizure disorder treatment options.
- Sec. 3. The state encourages state educational institutions that perform medical research and that are affiliated with a physician



1	registered under IC 16-42-28 to:
2	(1) conduct research by the physician; and
3	(2) participate in clinical studies or trials;
4	concerning the safety and efficacy of using cannabidiol in the
5	treatment of intractable epilepsy.
6	SECTION 12. IC 34-30-2-83.9 IS ADDED TO THE INDIANA
7	CODE AS A NEW SECTION TO READ AS FOLLOWS
8	[EFFECTIVE JULY 1, 2016]: Sec. 83.9. IC 16-42-28-11 (Concerning
9	the growing of industrial hemp and the manufacturing, processing,
10	suppling, and dispensing or use of cannabidiol in the treatment of
11	intractable epilepsy).
12	SECTION 13. IC 35-48-4-8.5, AS AMENDED BY P.L.208-2015,
13	SECTION 18, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
14	JULY 1, 2016]: Sec. 8.5. (a) A person who keeps for sale, offers for
15	sale, delivers, or finances the delivery of a raw material, an instrument,
16	a device, or other object that is intended to be or that is designed or
17	marketed to be used primarily for:
18	(1) ingesting, inhaling, or otherwise introducing into the human
19	body marijuana, hash oil, hashish, salvia, a synthetic drug, or a
20	controlled substance;
21	(2) testing the strength, effectiveness, or purity of marijuana, hash
22	oil, hashish, salvia, a synthetic drug, or a controlled substance;
23	(3) enhancing the effect of a controlled substance;
24	(4) manufacturing, compounding, converting, producing,
25	processing, or preparing marijuana, hash oil, hashish, salvia, a
26	synthetic drug, or a controlled substance;
27	(5) diluting or adulterating marijuana, hash oil, hashish, salvia, a
28	synthetic drug, or a controlled substance by individuals; or
29	(6) any purpose announced or described by the seller that is in
30	violation of this chapter;
31	commits a Class A infraction for dealing in paraphernalia.
32	(b) A person who knowingly or intentionally violates subsection (a)
33	commits a Class A misdemeanor. However, the offense is a Level 6
34	felony if the person has a prior unrelated judgment or conviction under
35	this section.
36	(c) This section does not apply to the following:
37	(1) Items marketed for use in the preparation, compounding,
38	packaging, labeling, or other use of marijuana, hash oil, hashish,
39	salvia, a synthetic drug, or a controlled substance as an incident
40	to lawful research, teaching, or chemical analysis and not for sale.
41	(2) Items marketed for or historically and customarily used in
42	connection with the planting, propagating, cultivating, growing,



1	harvesting, manufacturing, compounding, converting, producing
2	processing, preparing, testing, analyzing, packaging, repackaging
3	storing, containing, concealing, injecting, ingesting, or inhaling
4	of tobacco or any other lawful substance.
5	(3) A qualified entity (as defined in IC 16-41-7.5-3) that provides
6	a syringe or needle as part of a program under IC 16-41-7.5.
7	(4) Any entity or person that provides funding to a qualified entity
8	(as defined in IC 16-41-7.5-3) to operate a program described in
9	IC 16-41-7.5.
10	(5) A:
l 1	(A) licensed facility or manufacturer that manufacturers
12	processes, or supplies cannabidiol; or
13	(B) licensed grower who grows and supplies industria
14	hemp to a facility or manufacturer that manufacturers or
15	processes cannabidiol;
16	under IC 15-15-13.
17	(6) A physician described in IC 16-42-28 who:
18	(A) has been approved by the state department of health
19	under IC 16-42-28 to dispense or use cannabidiol in the
20	treatment of a patient for intractable epilepsy in a pilo
21	research study described in IC 16-42-28; and
22	(B) is dispensing or otherwise using the cannabidiol in the
23	treatment of the registered individual diagnosed with
24	intractable epilepsy.
25	(7) A physician, patient, or caregiver who is registered under
26	IC 16-42-28 for the use of cannabidiol in the treatment of
27	intractable epilepsy only if the cannabidiol is being used:
28	(A) for the patient; and
29	(B) in the manner;
30	approved by the state department of health for the purposes
31	of the registry.
32	SECTION 14. IC 35-48-4-10, AS AMENDED BY P.L.168-2014
33	SECTION 100, IS AMENDED TO READ AS FOLLOWS
34	[EFFECTIVE JULY 1, 2016]: Sec. 10. (a) A person who:
35	(1) knowingly or intentionally:
36	(A) manufactures;
37	(B) finances the manufacture of;
38	(C) delivers; or
39	(D) finances the delivery of;
10	marijuana, hash oil, hashish, or salvia, pure or adulterated; or
11	(2) possesses, with intent to:
12	(A) manufacture;



1	(B) finance the manufacture of;
2	(C) deliver; or
3	(D) finance the delivery of;
4	marijuana, hash oil, hashish, or salvia, pure or adulterated;
5	commits dealing in marijuana, hash oil, hashish, or salvia, a Class A
6	misdemeanor, except as provided in subsections (b) through (d).
7	(b) A person may be convicted of an offense under subsection (a)(2)
8	only if there is evidence in addition to the weight of the drug that the
9	person intended to manufacture, finance the manufacture of, deliver,
10	or finance the delivery of the drug.
11	(c) The offense is a Level 6 felony if:
12	(1) the person has a prior conviction for a drug offense and the
13	amount of the drug involved is:
14	(A) less than thirty (30) grams of marijuana; or
15	(B) less than five (5) grams of hash oil, hashish, or salvia; or
16	(2) the amount of the drug involved is:
17	(A) at least thirty (30) grams but less than ten (10) pounds of
18	marijuana; or
19	(B) at least five (5) grams but less than three hundred (300)
20	grams of hash oil, hashish, or salvia.
21	(d) The offense is a Level 5 felony if:
22 23 24 25	(1) the person has a prior conviction for a drug dealing offense
23	and the amount of the drug involved is:
24	(A) at least thirty (30) grams but less than ten (10) pounds of
	marijuana; or
26	(B) at least five (5) grams but less than three hundred (300)
27	grams of hash oil, hashish, or salvia; or
28	(2) the:
29	(A) amount of the drug involved is:
30	(i) at least ten (10) pounds of marijuana; or
31	(ii) at least three hundred (300) grams of hash oil, hashish
32	or salvia; or
33	(B) offense involved a sale to a minor.
34	(e) This section does not apply to the following:
35	(1) A:
36	(A) licensed facility or manufacturer that manufacturers
37	processes, or supplies cannabidiol; or
38	(B) licensed grower who grows and supplies industrial
39	hemp to a facility or manufacturer that manufacturers or
40	processes cannabidiol;
41	under IC 15-15-13.
42	(2) A physician described in IC 16-42-28 who:



1	(A) has been approved by the state department of health
2	under IC 16-42-28 to dispense or use cannabidiol in the
3	treatment of a patient for intractable epilepsy in a pilo
4	research study described in IC 16-42-28; and
5	(B) is dispensing or otherwise using the cannabidiol in the
6	treatment of the registered patient diagnosed with
7	intractable epilepsy.
8	(3) A physician, patient, or caregiver who is registered under
9	IC 16-42-28 for the use of cannabidiol in the treatment of
10	intractable epilepsy only if the cannabidiol is being used:
l 1	(A) for the patient; and
12	(B) in the manner;
13	approved by the state department of health for the purposes
14	of the registry.
15	SECTION 15. IC 35-48-4-11, AS AMENDED BY
16	P.L.226-2014(ts), SECTION 12, IS AMENDED TO READ AS
17	FOLLOWS [EFFECTIVE JULY 1, 2016]: Sec. 11. (a) A person who
18	(1) knowingly or intentionally possesses (pure or adulterated)
19	marijuana, hash oil, hashish, or salvia;
20	(2) knowingly or intentionally grows or cultivates marijuana; or
21	(3) knowing that marijuana is growing on the person's premises
22	fails to destroy the marijuana plants;
23	commits possession of marijuana, hash oil, hashish, or salvia, a Class
24	B misdemeanor, except as provided in subsections (b) through (c).
25	(b) The offense described in subsection (a) is a Class A
26	misdemeanor if the person has a prior conviction for a drug offense.
27	(c) The offense described in subsection (a) is a Level 6 felony if:
28	(1) the person has a prior conviction for a drug offense; and
29	(2) the person possesses:
30	(A) at least thirty (30) grams of marijuana; or
31	(B) at least five (5) grams of hash oil, hashish, or salvia.
32	(d) This section does not apply to the following:
33	(1) A:
34	(A) licensed facility or manufacturer that manufacturers
35	processes, or supplies cannabidiol; or
36	(B) licensed grower who grows and supplies industria
37	hemp to a facility or manufacturer that manufacturers or
38	processes cannabidiol;
39	under IC 15-15-13.
10	(2) A physician described in IC 16-42-28 who:
‡1 ‡2	(A) has been approved by the state department of health
L)	under IC 16-42-28 to dispense or use cannabidial in the



1	treatment of a patient for intractable epilepsy in a pilot
2	research study described in IC 16-42-28; and
3	(B) is dispensing or otherwise using the cannabidiol in the
4	treatment of the registered patient diagnosed with
5	intractable epilepsy.
6	(3) A physician, patient, or caregiver who is validly registered
7	under IC 16-42-28 for the use of cannabidiol in the treatment
8	of intractable epilepsy only if the cannabidiol is being used:
9	(A) for the patient; and
10	(B) in the manner;
11	approved by the state department of health for the purposes
12	of the registry.

